

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/08/25 18:36:27

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR3084867.sorted.bam -c -nw 400 -hm 3
```

1.2. Alignment

Command line:	/home/luna/Desktop/Software/bwa/bwa mem -t 16 -k 30 -R @RG\tID:Singlecell2022\tLB:Library\tPL:ILLUMINA\tSM:sample_SRR3084867 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR3084867.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Sun Aug 25 18:36:26 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR3084867.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	2,705,038
Mapped reads	2,421,321 / 89.51%
Unmapped reads	283,717 / 10.49%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	18,403 / 0.68%
Read min/max/mean length	30 / 76 / 76.24
Duplicated reads (estimated)	106,697 / 3.94%
Duplication rate	3.69%
Clipped reads	801,678 / 29.64%

2.2. ACGT Content

Number/percentage of A's	49,350,000 / 29.34%
Number/percentage of C's	30,765,382 / 18.29%
Number/percentage of T's	53,969,701 / 32.09%
Number/percentage of G's	34,063,387 / 20.25%
Number/percentage of N's	26,284 / 0.02%
GC Percentage	38.55%

2.3. Coverage

Mean	0.0543

Standard Deviation	0.4108
--------------------	--------

2.4. Mapping Quality

Mean Mapping Quality	47.68
----------------------	-------

2.5. Mismatches and indels

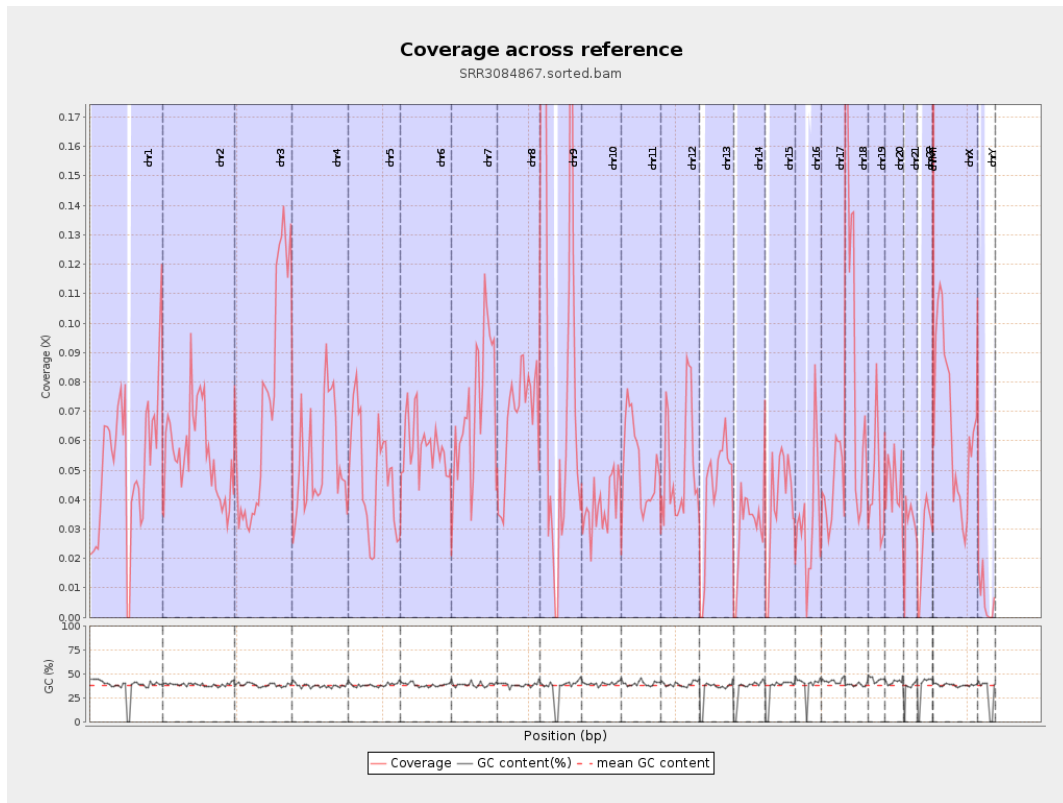
General error rate	0.84%
Mismatches	1,386,869
Insertions	13,242
Mapped reads with at least one insertion	0.54%
Deletions	36,394
Mapped reads with at least one deletion	1.49%
Homopolymer indels	48.33%

2.6. Chromosome stats

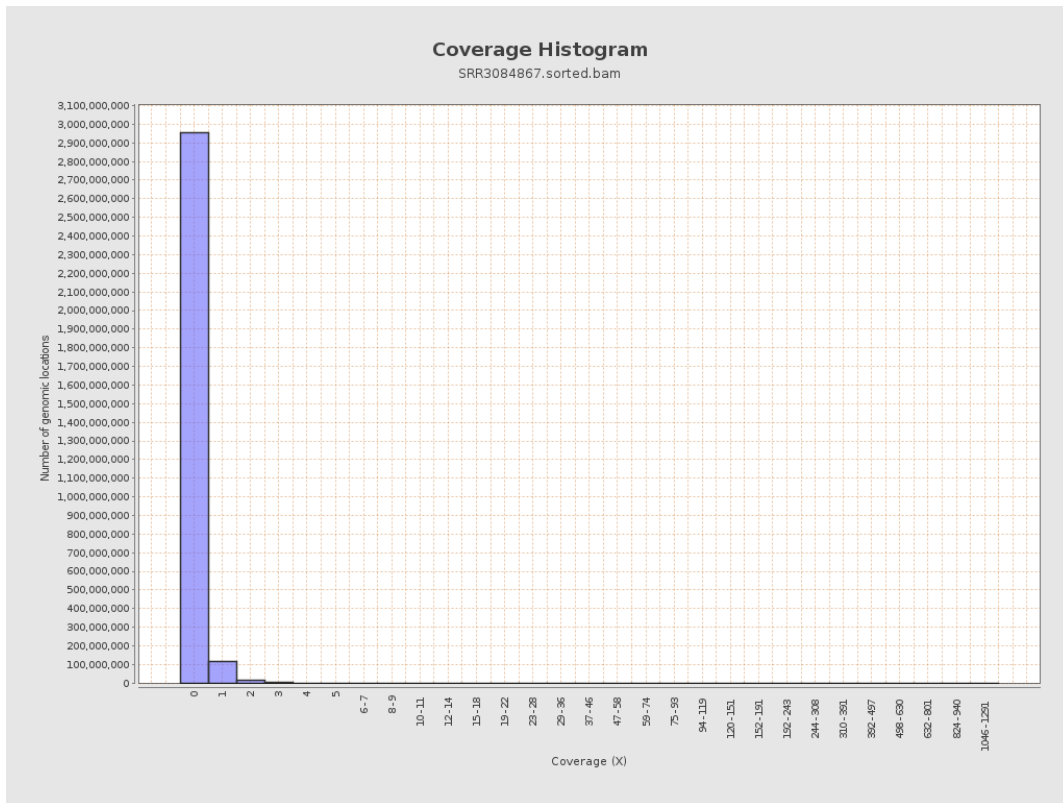
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	13036334	0.0523	0.6434
chr2	243199373	13518322	0.0556	0.4299
chr3	198022430	14290663	0.0722	0.3097
chr4	191154276	10221488	0.0535	0.2687
chr5	180915260	8696534	0.0481	0.2575
chr6	171115067	9970829	0.0583	0.2864
chr7	159138663	11252630	0.0707	0.4532

chr8	146364022	9872748	0.0675	0.8801
chr9	141213431	13152689	0.0931	0.4677
chr10	135534747	5156103	0.038	0.2964
chr11	135006516	7017156	0.052	0.3427
chr12	133851895	6888387	0.0515	0.2632
chr13	115169878	5000520	0.0434	0.2374
chr14	107349540	3239616	0.0302	0.2113
chr15	102531392	3944198	0.0385	0.2238
chr16	90354753	2856731	0.0316	0.2284
chr17	81195210	3690256	0.0454	0.2535
chr18	78077248	7229576	0.0926	0.6908
chr19	59128983	2740654	0.0464	0.4148
chr20	63025520	2885079	0.0458	0.25
chr21	48129895	1502511	0.0312	0.2171
chr22	51304566	1319633	0.0257	0.1795
chrMT	16571	66169	3.9931	2.7779
chrX	155270560	10309372	0.0664	0.3199
chrY	59373566	381224	0.0064	0.2063

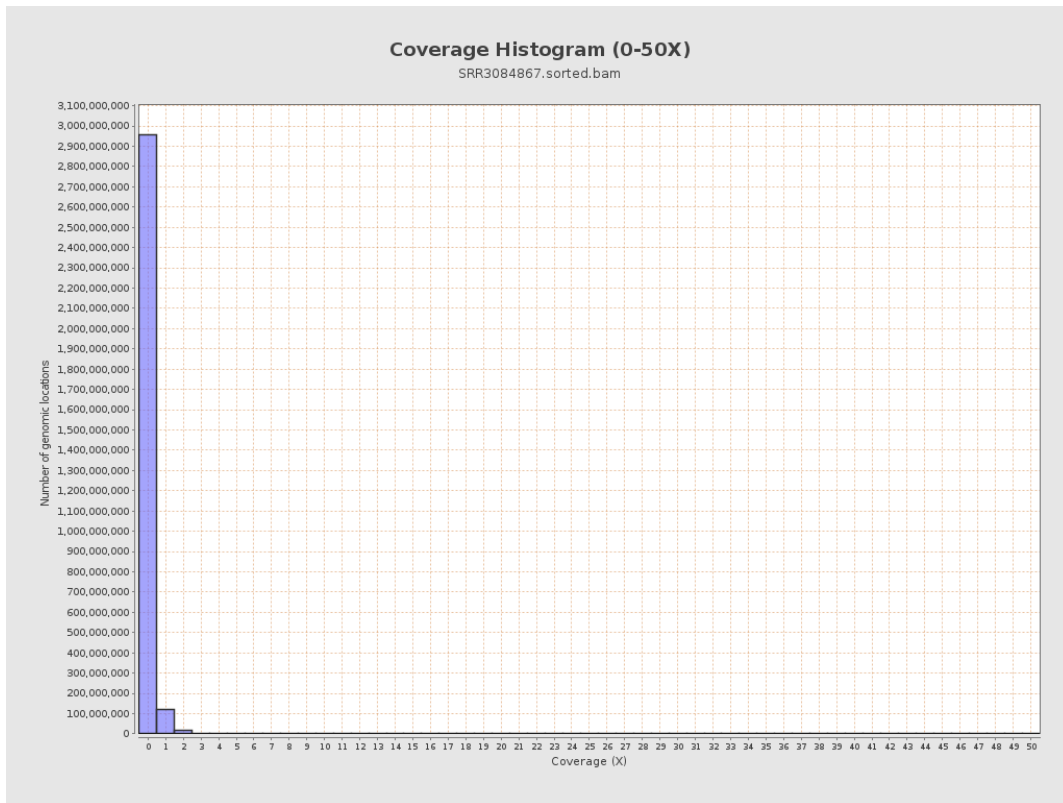
3. Results : Coverage across reference



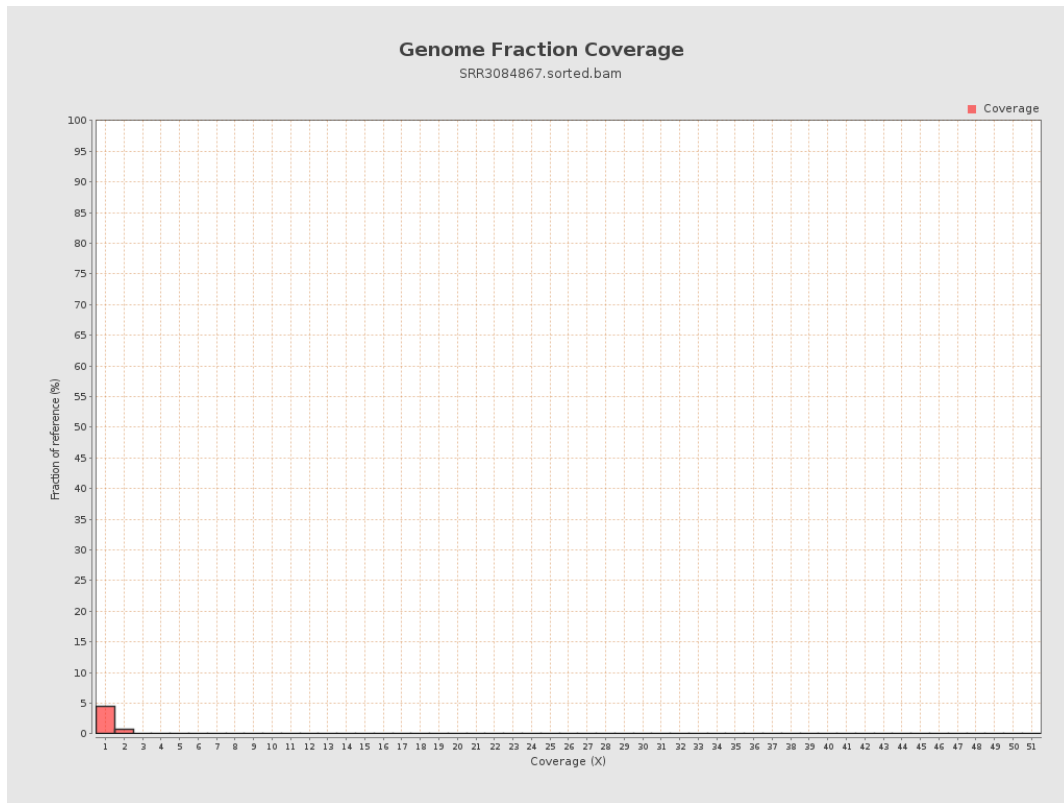
4. Results : Coverage Histogram



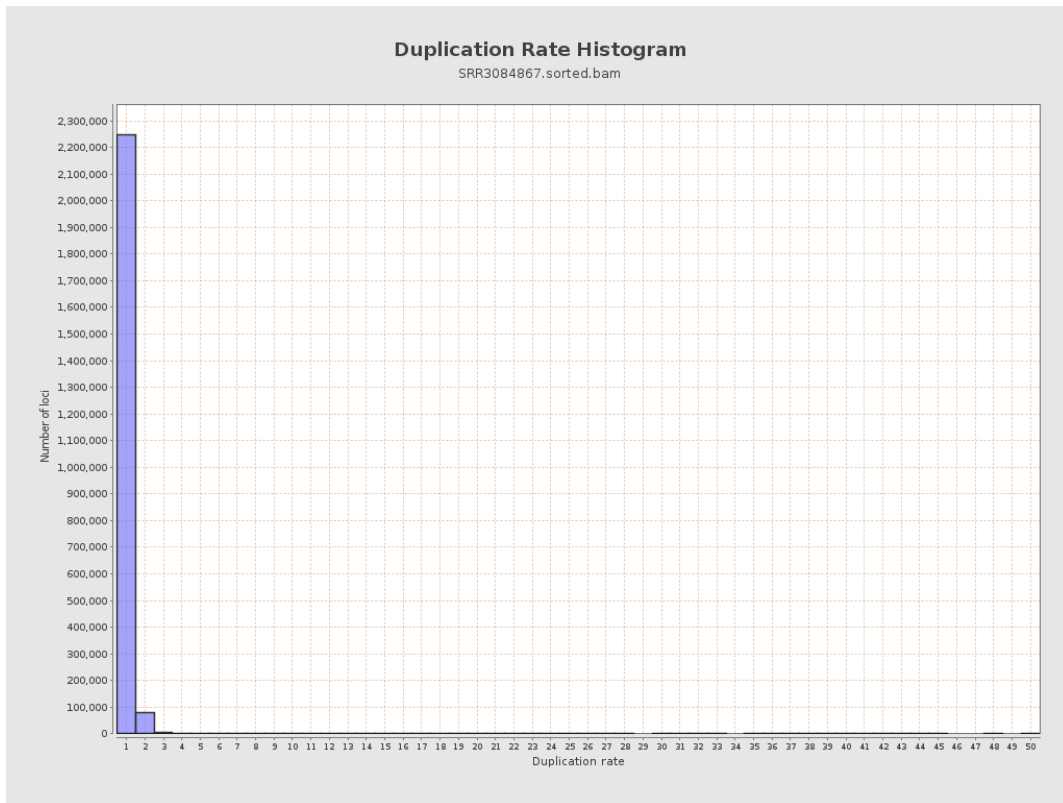
5. Results : Coverage Histogram (0-50X)



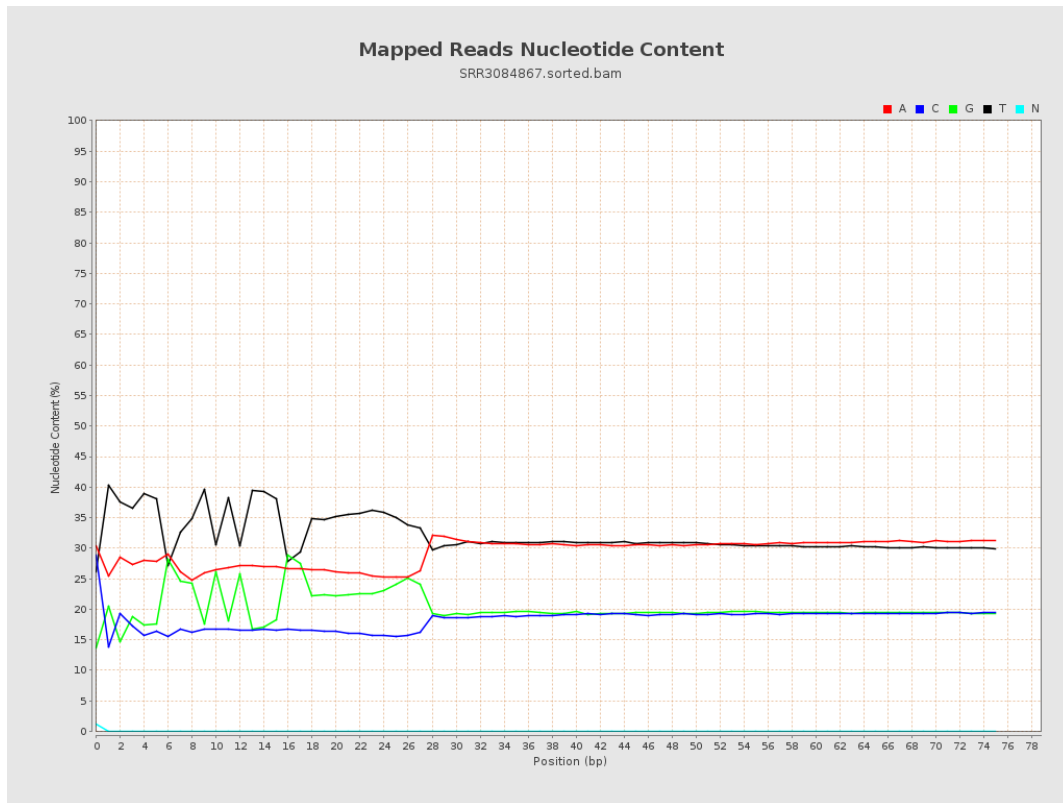
6. Results : Genome Fraction Coverage



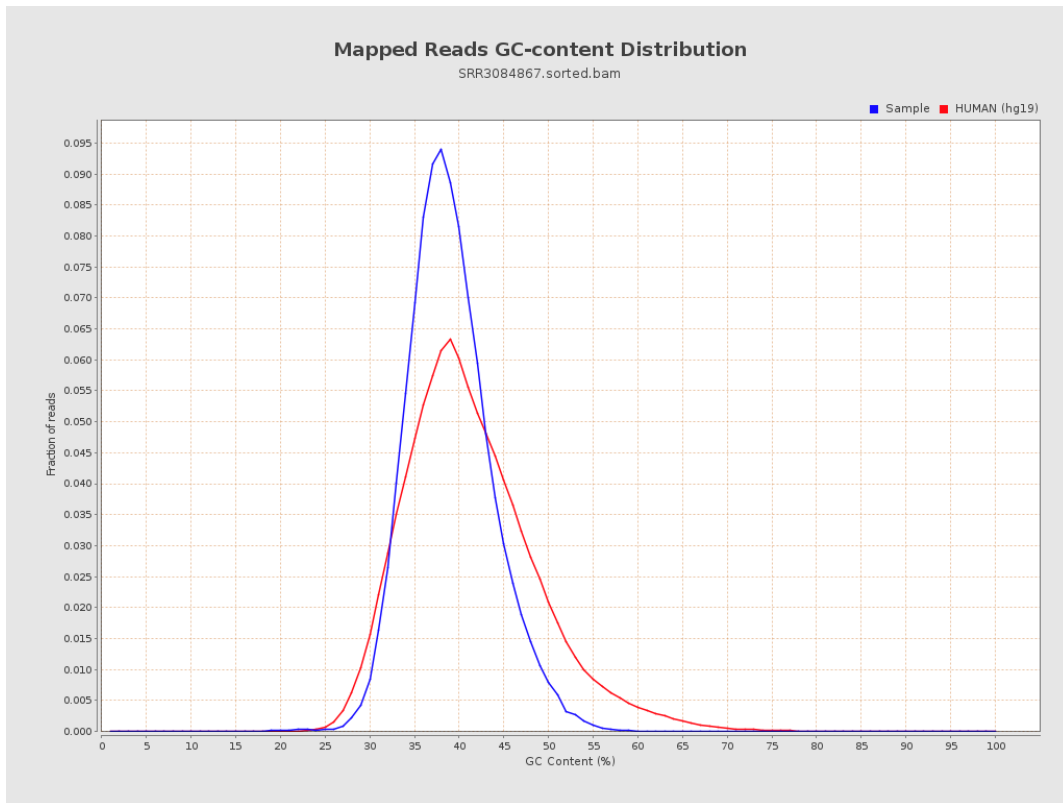
7. Results : Duplication Rate Histogram



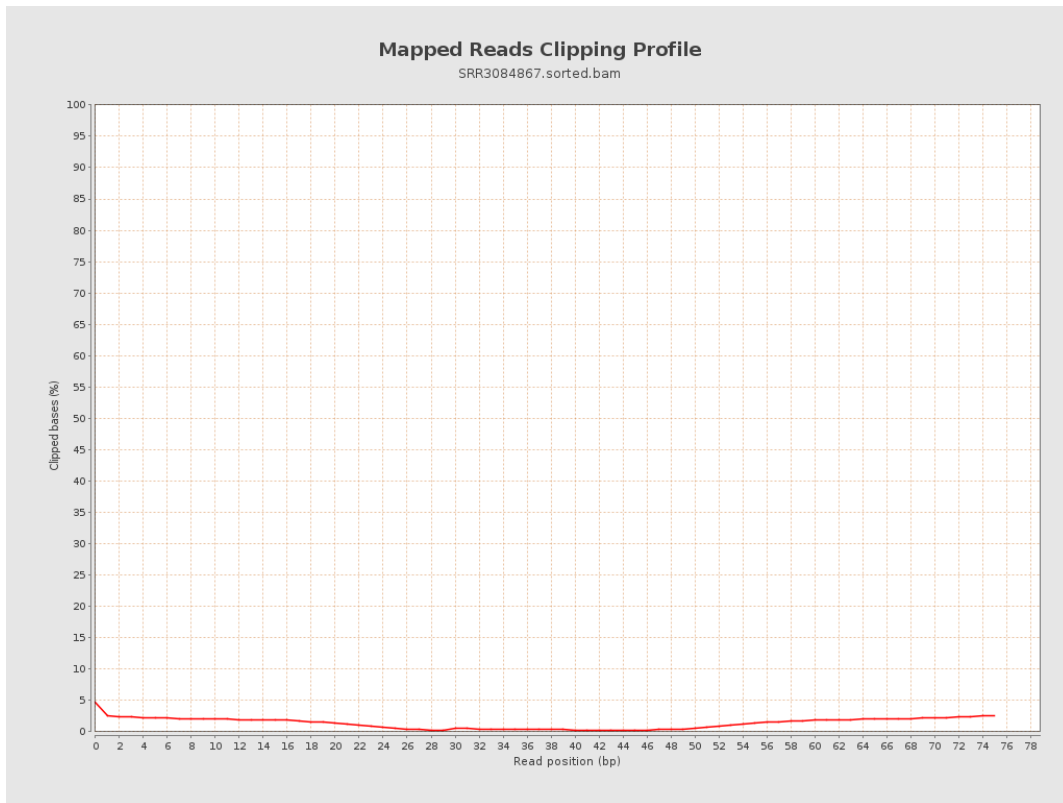
8. Results : Mapped Reads Nucleotide Content



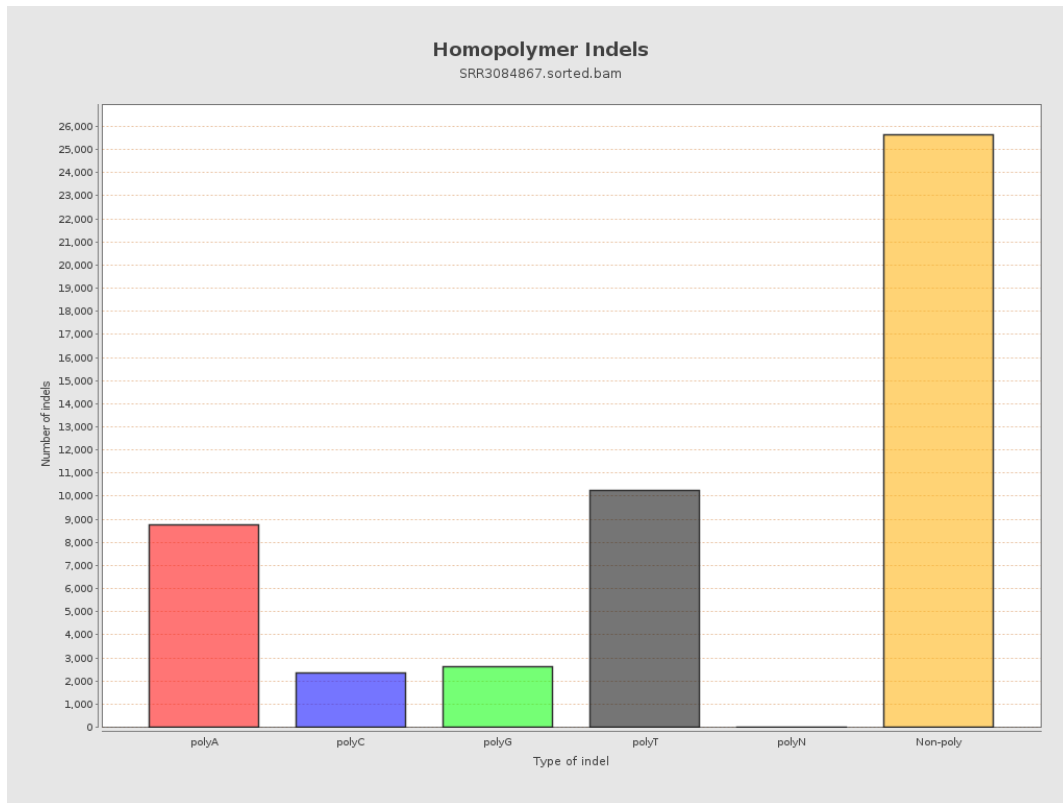
9. Results : Mapped Reads GC-content Distribution



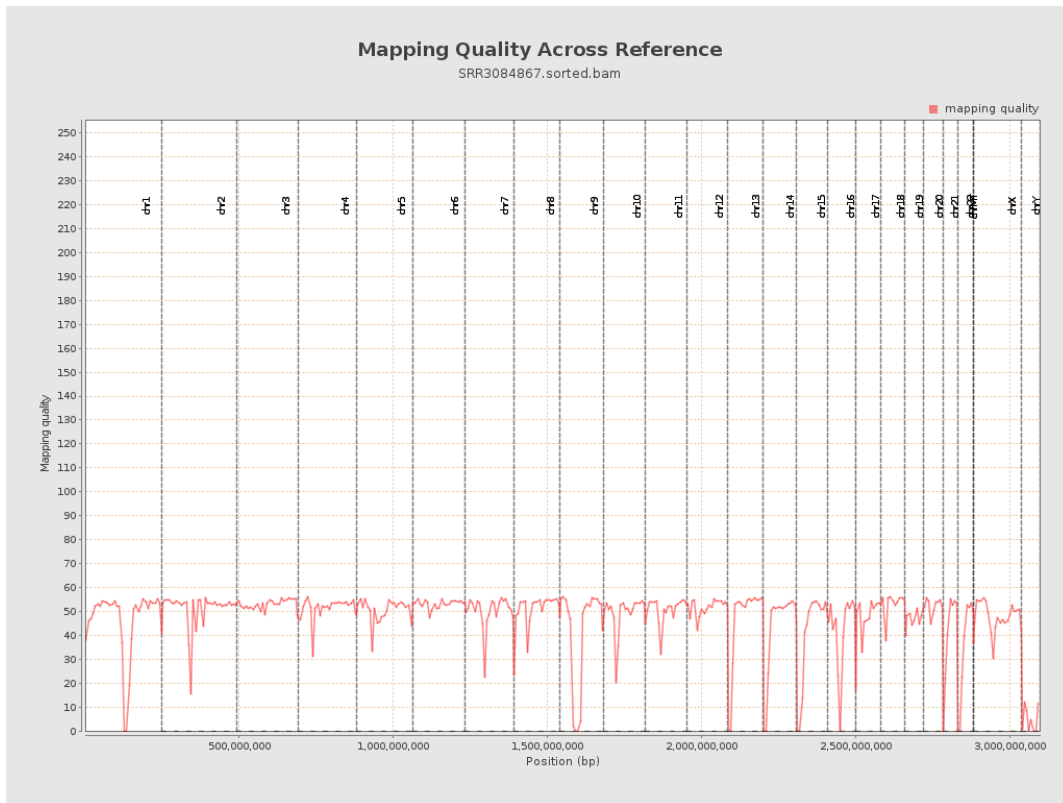
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

